

K150712 LTM-Perforated Surgical MeshJul 2, 2015
105 days to decisionK150712 · Product code: **FTM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k150712/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical (FTM)
Date received	Mar 19, 2015
Decision date	Jul 2, 2015
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lifecell Corporation
Location	Branchburg, NJ, US
Contact	Linda Scamardella
510(k) history	5 submissions · 5 cleared · 2014-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k150712/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 20, 2026